ULTRA CLEAN ANTISEPTIC HAND SANITIZER- alcohol gel
O.C.C.S. Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

ULTRA CLEAN ANTISEPTIC HAND SANITIZER

Drug Facts

Active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Uses

• hand sanitizer to decrease bacteria on the skin
• recommended for repeated use
• for use when soap and water are not available

Warnings

Flammable, keep away from fire/flame
For external use only

Do not use
• in children less than 2 months of age
• on open skin wounds

When using this product • do not get into eyes. In case of contact, rinse eyes thoroughly with water

Stop use and ask a doctor if
• irritation and redness develop
• condition persists for more than 72 hours

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

• wet hands thoroughly with product and allow to dry without wiping
• supervise children under 6 years of age when using this product to avoid swallowing

Other information

• store between 15-30°C (59-86°F)
• avoid freezing and excessive heat above 40°C (104°F)

Inactive ingredients

glycerin, carbomer, aminomethylpropanol, water

***Questions?* +1-714-816-3040**

You may also report serious side effects to this phone number.
Mon-Fri 9:00 AM - 5:00 PM

The Professional Line of Chemicals

70% Non-Scented Alcohol

Kills More Than 99.9% of Germs

Leaves Hands Silky Smooth

Reduces bacteria on Hands

Description

This is a premium alcohol based hand sanitizer. It reduces bacteria on hands while leaving the skin silky soft.

Manufactured by:

OCCS INC.

10680 Fern Ave.

Stanton, CA 90680

USA

(714) 816-3040

www.occsinc.com

Packaging

ULTRA CLEAN ANTISEPTIC HAND SANITIZER

alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78263-326
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 L in 100 L

Inactive Ingredients	
Ingredient Name	**Strength**
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78263-326-41	3.78 L in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020	08/20/2023
2	NDC:78263-326-43	0.946 L in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020	08/20/2023
3	NDC:78263-326-45	0.236 L in 1 BOTTLE; Type 0: Not a Combination Product	06/01/2020	08/20/2023

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	06/01/2020	08/20/2023

Labeler - O.C.C.S. Inc. (012261504)

Establishment

Name	Address	ID/FEI	Business Operations
O.C.C.S. Inc.		012261504	manufacture(78263-326)

Revised: 8/2021 O.C.C.S. Inc.